



Drug Enforcement Administration

[Docket No. DEA-838]

Importer of Controlled Substances Application: SpecGX, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: SpecGX, LLC. has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before **[INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER]**. Such persons may also file a written request for a hearing on the application on or before **[INSERT DATE 30 DAYS AFTER DATE OF PUBLICATION IN THE FEDERAL REGISTER]**.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on February 5, 2021, SpecGX LLC, 3600 North 2nd Street, Saint Louis, Missouri

63147, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled Substance	Drug Code	Schedule
Marihuana	7360	I
Phenylacetone	8501	II
Coca Leaves	9040	II
Thebaine	9333	II
Opium, Raw	9600	II
Poppy Straw Concentrate	9670	II
Tapentadol	9780	II

The company plans to import the listed controlled substances for bulk manufacture into Active Pharmaceutical Ingredients (API) for distribution to its customers. In reference to Tapentadol (9780) and Thebaine (9333), the company plans to import intermediate forms of these controlled substances for further manufacturing prior to distribution to its customers. In reference to drug code 7360 (Marihuana), the company plans to import synthetic cannabinol. No other activity for this drug is authorized for this registration. Placement of these codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2).

Authorization will not extend the import of Food and Drug Administration-approved or non-approved finished forms for commercial sale.

William T. McDermott,

Assistant Administrator.

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